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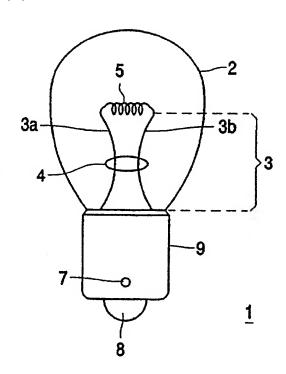
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(54) Title: RED-COLORED ELECTRIC LAMP



(57) Abstract: The invention relates to electric lamps and more particularly to automotive signal lamps. A glass composition is described for use in electric lamps, which glass is red colored. The glass composition comprises copper between 0.1 and 2% by weight, tin between 0.1 and 2% by weight, barium between 7 and 11% by weight, and strontium between 1 and 5% by weight.



F1, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)

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Red-colored electric lamp.

5 FIELD OF THE INVENTION

The present invention relates to electric lamps. In particular the invention relates to an electric lamp having a glass envelope, the glass being red colored.

The invention applies to incandescent lamps and more particularly to automotive signal lamps, e.g. indicator lamps.

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BACKGROUND OF THE INVENTION

Automotive signal lamps must have specific colors defined by international traffic regulations. For example, automotive signalling functions like tail, stop, or fog lighting require red lamps. In order to obtain the red color as defined by international traffic regulations, prior-art lamps are usually coated with a red varnish or lacquer. A disadvantage of the prior-art lamps is that the lacquer or varnish may degrade and may eventually peel off from the wall of the lamp envelope. Actually, these red coatings usually consist of vulnerable constituents that have a relative low thermal resistance. Another disadvantage of the prior-art lamps is that additional steps are necessary during the manufacturing process of the lamp for cleaning and varnishing the lamp envelope. The manufacturing cost is increased by these additional steps.

SUMMARY OF THE INVENTION

It is an object of the invention to provide an electric lamp suitable for automotive applications, said lamp having a glass envelope, the glass being red colored.

According to the invention, these objects are achieved by an electric lamp as mentioned in the opening paragraph, the glass composition comprising copper between 0.1 and 2% by weight, tin between 0.1 and 2% by weight, barium between 7 and 11% by weight, and strontium between 1 and 5% by weight.

No additional manufacturing step is necessary during the production of an electric lamp according to the invention, since the specific color point of the glass envelope is realized by the particular composition of the glass. At least, the steps of cleaning and varnishing are suppressed. Due to its specific composition, the 5

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glass may have a red color point suitable for automotive signal lamps, as defined by international traffic regulations.

Moreover, the colored glass obtained in accordance with this glass composition yields a better quality glass, since the obtained red color inherent in the glass composition stays forever, which may not be the case of varnished glasses. Actually, the use of barium and strontium lowers the ion conductivity of the glass, which results in a thermostable color of the glass at lamp operating temperatures.

Furthermore, such a glass is particularly suitable for automotive signal lamps, because the presence of barium increases the electrical resistivity of the glass, which is required for the electrical feed-through.

In an advantageous embodiment, the glass composition comprises the following constituents, expressed in percentages by weight:

SiO ₂	60-72
Al ₂ O ₃	1-5
Li ₂ O	0.5-1.5
Na ₂ O	5-9
K ₂ O	3-7
MgO	1-2
CaO	1-3
SrO	1-5
ВаО	7-11
CuO	0.1-2
SnO_2	0.1-2

15 The SiO₂ content is limited to 60-72%. These contents, in combination with the other constituents, lead to a readily meltable glass. Al₂O₃ improves the chemical resistance and the corrosion resistance of the glass. The alkaline metal oxides Li₂O, Na₂O, and K₂O are used as melting agents and reduce the viscosity of the glass. MgO and CaO reduce the liquefying temperature and the melting 20 temperature of the glass. Such a glass is lead-free, cadmium-free, and selenium-free, which elements are harmful for the environment.

The invention and additional features, which may be optionally used to implement the invention, are apparent from and will be elucidated with reference to the drawings described hereinafter.

5 BRIEF DESCRIPTION OF THE DRAWINGS

The invention will now be described in more detail, by way of example, with reference to the accompanying drawings, in which:

- Fig. 1 is a schematic chart illustrating an example of an electric lamp according to the invention.
- Fig. 2 is a chromaticity diagram showing the characteristics in the color triangle of red colored glasses according to the invention.

DETAILED DESCRIPTION OF THE INVENTION

- Fig. 1 shows an electric automotive signal lamp 1, also denoted indicator lamp, comprising:
 - a lamp envelope or bulb 2 made from red colored glass, having a wall thickness comprised between, for example, 0.3 mm and 1.1 mm,
 - a mount 3 comprising two lead wires 3a and 3b held by a bead 4 mounted inside the glass bulb 2 for supporting a coiled filament 5,
 - an exhaust tube 7, heated and sealed up with the mount 3 and the bulb 2, thereby forming a vacuum-tight pinch for exhausting gases from the lamp envelope 2 and for introducing inert gas into the lamp envelope during the manufacturing process of the lamp,
 - an electric contact 8 connected to the mount 3 for the lamp to be electrically supplied by the electric circuits of the car,
 - a metal cap 9 to fit in a holder of the lamp set.

In an advantageous embodiment of the invention, the starting materials for making the glass are quartz sand, spodumene, dolomite, the carbonates of Li, Na, K, Sr, and Ba, CuO and SnO₂.

An electric lamp in accordance with an advantageous embodiment of the invention has a glass bulb with a glass composition comprising the constituents as listed in Table 1.

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Constituent	Composition
	(% by weight)
SiO ₂	60-72
Al ₂ O ₃	1-5
Li ₂ O	0.5-1.5
Na ₂ O	5-9
K ₂ O	3-7
MgO	1-2
CaO	1-3
SrO	1-5
BaO	7-11
CuO	0.1-2
SnO_2	0.1-2

Table 1: glass composition according to an advantageous embodiment of the invention.

In this table, the percentages by weight of the oxidic forms of the constituents are given. This does not necessary imply that these elements are present in the glass in this oxidic form. For example, copper in the glass might be present as metallic Cu, Cu⁺, CuO, or other forms of copper. As a consequence, expressions like "the glass composition comprises element X between α and β % by weight" mean that all the forms of element X comprised in the glass composition account for a percentage between α and β % by weight.

SiO₂ serves as a network former in the glass. The SiO₂ content is limited to 60-72% by weight, leading to a readily meltable glass in combination with the other constituents. If the content is below 60% by weight, the cohesion of the glass and the chemical resistance are reduced. If the content is above 72% by weight, the glass formation is impeded and the risk of surface crystallization increases. Al₂O₃ improves the chemical resistance and the corrosion resistance of the glass. Below 1% by weight the effect is too small and the crystallization tendency of the glass increases. Above 5% by weight the viscosity and the softening temperature of the glass increase too much, which affects the workability of the glass. The

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alkaline metal oxides Li₂O, Na₂O, and K₂O are used as melting agents and reduce the viscosity of the glass. They increase the electrical resistance of the glass (mixed-alkali effect). BaO has the favorable property that it increases the electrical resistance of the glass and reduces the softening temperature of the glass. Below 7% by weight, the melting temperature, the softening temperature, and the working temperature increase too much. Above 11% by weight, the liquidus temperature and hence the crystallization tendency increase too much. The alkaline-earth metal oxides SrO, MgO, and CaO have the favorable property that they reduce the liquefying temperature and the melting temperature of the glass.

The glass may additionally contain some Fe₂O₃ as an impurity originating from the raw materials used. Also TiO₂, ZrO₂, and MnO may be found as trace elements.

The red color of the glass according to the invention is obtained by adding between 0.1 and 2% by weight of CuO and a reducing agent, e.g. carbone or charcoal, in the glass composition. SnO₂ is also added in the glass composition.

In the presence of a reducing agent and SnO₂, copper colors silicate glasses red, which is explained by the possible precipitation of Cu in metallic form, as will be described hereinafter.

An electric lamp in accordance with another embodiment of the invention has a glass bulb with a glass composition comprising the constituents as listed in Table 2.

Constituent	Composition
	(% by weight)
SiO ₂	60-72
$\mathrm{Al_2O_3}$	1-5
Li ₂ O	0.5-1.5
Na ₂ O	5-9
K ₂ O	·3-7
MgO	1-2
CaO	1-3
SrO	1-5
BaO	7-11

CuO	0.6-0.9		
SnO_2	0.3-1.8		
Fe ₂ O ₃	0 - 0.05		

Table 2: glass composition according to another embodiment of the invention.

With respect to the element whose weight percentage has a lower limit equal to zero, it means that the element must not be added as a raw material may but yet be present in the finished glass as a result of contamination of the raw materials.

The glasses made with the constituents as listed in Table 1 or 2 have characteristics as shown in Table 3.

Physical property	Value		
T _{strain} (low stress relief temperature)	455°C		
T _{anneal} (high stress relief temperature)	490°C		
T _{soft} (softening temperature)	675°C		
T _{melt} (melting temperature)	1490°C		
Specific mass	$2.62*10^3$ kg.m ³		
Specific resistance	$7.9*10^6\Omega$.m		
Linear expansion coefficient (25-300°C)	9.1*10 ⁻⁶ /°C		

Table 3: Physical properties of the glass according to the invention.

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The glass-making process will now be described below. A batch is prepared by weighing and mixing all oxidic components mentioned above. The batch is fed to a conventional continuous melting tank. Adjustments of the recipe can be made by adding separate ingredients in the feeding system for enhancing the coloring. The melting conditions, including temperature and gas atmosphere, are adjusted to obtain a stable processing for melting, refining, and color stabilization.

In a series of complex reactions, the excess gases, including CO, are removed from the glass melt to avoid air lines in the formed tubes. Glass tubes are made at the end of the glass manufacturing process using the well-known Danner WO 2004/041739 PCT/IB2003/004580

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process in the factory. However, the well-known Vello process could also be applied. Part of the tube is heated to make lamp bulbs.

The striking of the glass may be explained by the presence of metallic Cu in crystals of colloidal size, in equilibrium with Cu⁺. Furthermore, it appears that striking requires a certain degree of nucleation, which is why SnO₂ is added. Cu is introduced in the form of the oxide in combination with a reducing agent. In this example, the reducing agent is carbon, but any other reducing agent could be used, including Al and Si. Suitable time and temperature of striking, which depend on the basic composition, are best determined in a gradient furnace. For example, a striking temperature between 500 and 550°C might be used. The time of striking might be between 0.5 and 3 hours.

Copper is introduced as CuO in a percentage comprised between 0.1 and 2 % by weight. Larger amounts of copper are harmful, as they decrease transmissivity in the red region. Lower concentrations do not produce coloring. Under reducing conditions, an equilibrium is obtained between Cu^{2+} , Cu^{+} and Cu. When sufficient Cu^{+} ions are present in the glass, the concentration of Cu in equilibrium with Cu^{+} exceeds the solubility limits and Cu is precipitated. The red color is obtained by colloidally dispersed Cu particules in equilibrium with Cu^{+} , when the bivalent copper has disappeared. The reducing element should therefore be able to reduce all Cu^{2+} . In this example, the reducing element C is chosen in a proportion C/Cu = 1.67.

The function of tin may be that of a metallophilic element. As soon as a nucleus of a Cu crystal has formed, it selectively attracts the Sn²⁺ ions, which form an efficient barrier against further growth of the metal crystal and thus stabilizes the colloidal dispersion of the Cu and prevents further growth of the Cu atom aggregates and their overstriking.

Fig. 2 shows a variety of color points of red colored glass for use in the glass envelope of signal lamps in accordance with international traffic regulations, with respect to X and Y co-ordinates in the chromaticity chart. This color point may be achieved with the glass composition in accordance with the invention described above. It is noted that the exact composition of the glass which gives the color point

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WO 2004/041739 PCT/IB2003/004580

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depends on the production of the glass, particularly the reducing state, and on the curing after bulb blowing.

In Europe, a red color point for automotive indicator lamps is defined by the ECE regulation, known to those skilled in the art. It corresponds to the area shown in continuous lines of Fig. 2. The red ECE-region is defined by the following color coordinates: (0.657, 0.335); (0.665, 0.335); (0.732, 0.27); (0.728, 0.27).

The SAE (Society of Automotive Engineers) has proposed another region, which is defined by the following color coordinates: (0.65, 0.33); (0.67, 0.33); (0.7367, 0.2653); (0.7164, 0.2636). This region is shown in dotted lines on Fig. 2.

The GTB (Groupe de Travail de Bruxelles) commission has proposed a new, larger area which encloses both the current ECE and SAE color boundaries. This is known as the CIE-region.

The triangles in the chart of Fig.2 show the color points obtained with a glass of 0.5 mm thickness comprising 0.9% by weight of copper, 0.6% by weight of tin, 9.0% by weight of barium, and 3.0% by weight of strontium. The striking temperature is 540°C. The results are given for striking times of 2 and 3 hours.

The drawings and their description hereinbefore illustrate rather than limit the invention. It will be evident that there are numerous alternatives which fall within the scope of the appended claims. In this respect, the following closing remark is made.

The word "comprising" does not exclude the presence of other elements than those listed in a claim. The word "a" or "an" preceding an element does not exclude the presence of a plurality of such elements.

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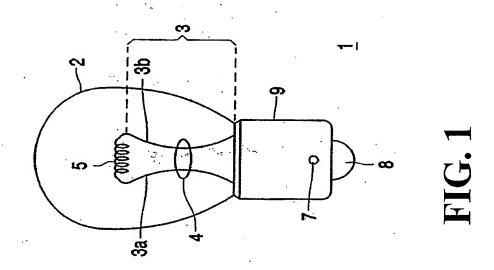
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<u>Claims</u>

- 1. An electric lamp having a glass envelope, the glass being red colored, the glass composition comprising copper between 0.1 and 2% by weight, tin between 0.1 and 2% by weight, barium between 7 and 11% by weight, and strontium between 1 and 5% by weight.
- 2. An electric lamp as claimed in claim 1, wherein the glass composition, expressed in percents by weight of the oxidic forms of the constituents, comprises:

SiO_2	60-72
Al ₂ O ₃	1-5
Li ₂ O	0.5-1.5
Na ₂ O	5-9
K ₂ O	3-7
MgO	1-2
CaO	1-3
SrO	1-5
BaO	7-11
CuO	0.1-2
SnO_2	0.1-2



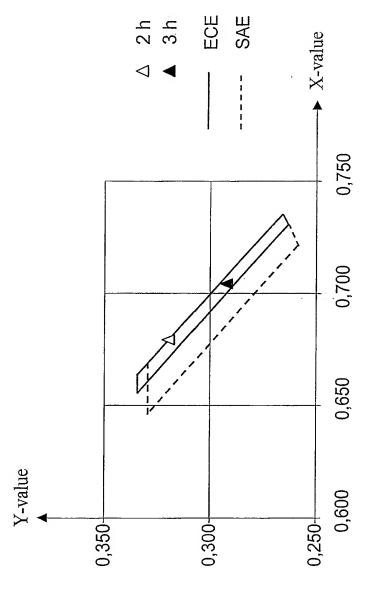


FIG. 2

INTERNATIONAL SEARCH REPORT

pcT/IB 03/04580

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C03C3/097 C03C4/02 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) IPC 7 C03C Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, CHEM ABS Data C. DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. Υ WO 02 46116 A (KONINKL PHILIPS ELECTRONICS 1,2 NV) 13 June 2002 (2002-06-13) page 3, line 7 -page 4, line 15 page 1, line 11 -page 2, line 11 Υ EP 0 603 933 A (PHILIPS ELECTRONICS NV) 1,2 29 June 1994 (1994-06-29) page 2, line 39 -page 3, line 36 Υ DE 100 53 450 A (SCHOTT DESAG AG) 1,2 8 May 2002 (2002-05-08) paragraphs '0014!-'0022!,'0026!; examples Further documents are listed in the continuation of box C. χ Patent family members are listed in annex. Special categories of cited documents: "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled in the art. other means "P" document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 9 January 2004 22/01/2004 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016 Van Bommel, L

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INTERNATIONAL SEARCH REPORT

Inter Application No
PCT/IB 03/04580

		PUI/IB U.	5/ 04300
	ation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No.
Y	DATABASE WPI Section Ch, Week 199614 Derwent Publications Ltd., London, GB; Class L01, AN 1996-131170 XP002266408 & ES 2 081 249 A (CRISTALERIAS MATARO SCOCL), 16 February 1996 (1996-02-16) abstract		1,2
Y	DATABASE WPI Section Ch, Derwent Publications Ltd., London, GB; Class L01, AN 1973-19338U XP002266409 & JP 48 010047 B (MITSUBISHI ELECTRIC CORP) abstract		1,2
			•
			•••
			·

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INTERNATIONAL SEARCH REPORT

Interr Application No
PCT/IB 03/04580

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 0246116	Α	13-06-2002	CN	1396891 T	12-02-2003
			WO	0246116 A1	13-06-2002
			EP	1345861 A1	24-09-2003
			US	2002117950 A1	29-08-2002
EP 0603933	A	29-06-1994	EP	0603933 A1	29-06-1994
			ΑU	5244993 A	23-06-1994
			BR	9305027 A	21-06-1994
			CA	2111145 A1	15-06-1994
			CN	1088895 A ,B	06-07-1994
			DE	69305311 D1	14-11-1996
			DE	69305311 T2	24-04-1997
			ES	2094998 T3	01-02-1997
			HU	69747 A2	28-09-1995
			JP	3299615 B2	08-07-2002
			JP	6206737 A	26-07-1994
			MX	9307816 A1	31-08-1994
			PL	301410 A1	27-06-1994
			SG	44912 A1	19-12-1997
			US	5470805 A	28-11-1995
DE 10053450	Α	08-05-2002	DE	10053450 A1	08-05-2002
ES 2081249	Α	16-02-1996	ES	2081249 A1	16-02-1996
JP 48010047	В		NONE		